

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

PATENT NO. : 7,981,601 B2
APPLICATION NO. : 10/566697
DATED : July 19, 2011
INVENTOR(S) : Heng Wang and Qiliang Cai

Page 1 of 4

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

Coversheet, FOREIGN PATENT DOCUMENTS

Foreign Patent Document reads “EP	0198328	A1	10/1986”
should read -- EP	0198328	A2	10/1986 --

Abstract, line 8, reads “tion results in the induction of high level of specific antibodies”
should read -- tion results in the induction of a high level of specific antibodies --

Column 1, line 44, reads “Ther, 3 (1), 31-36 (2001)), there is no related literature or”
should read -- Ther, 3 (1), 31-36 (2001)), there are no related publications or --

Column 1, line 66, reads “therapy of allergic response and tolerance in new born infants.”
should read -- therapy of allergic response and tolerance in newborn infants. --

Column 2, line 13, reads “binant vaccines. Moreover, it is problem that the synthesis of”
should read -- binant vaccines. Moreover, it is a problem that the synthesis of --

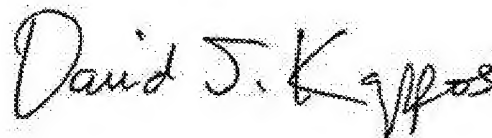
Column 2, line 60, reads “vaccines are directed can not generate satisfactory protective”
should read -- vaccines are directed cannot generate satisfactory protective --

Column 3, line 35, reads “according to the results of step e) and f);”
should read -- according to the results of steps e) and f); --

Column 3, line 42, reads “a) selecting, synthesizing and cloning into a vector a plu-”
should read -- a) selecting, synthesizing and cloning into a vector plu- --

Column 4, line 16, reads “ture of random assembled bi-epitope genes as templates (in”
should read -- ture of randomly assembled bi-epitope genes as templates (in --

Signed and Sealed this
Twentieth Day of December, 2011



David J. Kappos
Director of the United States Patent and Trademark Office

U.S. Pat. No. 7,981,601 B2

Column 4, line 18, reads “respectively), and in a 50 ul system primer free polymerase”
should read -- respectively), and in a 50 ul system a primer-free polymerase --

Column 5, line 4, reads “A. A blood smear of Plasmodium falciparum 3D7; B. A”
should read -- A. A blood smear of Plasmodium falciparum 3D7; B. A --

Column 5, line 5, reads “blood smear of Plasmodium falciparum FCC1; C. A blood”
should read -- blood smear of Plasmodium falciparum FCC1; C. A blood --

Column 6, line 31, reads “immunogenic in the literature on Plasmodium falciparum”
should read -- immunogenic in the literature on Plasmodium falciparum --

Column 6, line 52, reads “which differ from that with cDNA expression libraries, lie in”
should read -- which differ from that with cDNA expression libraries, lie in the fact --

Column 7, line 18, reads “In order to inhibit the growth of Plasmodium falciparum”
should read -- In order to inhibit the growth of Plasmodium falciparum --

Column 7, line 23, reads “stages of Plasmodium falciparum which are homologous to”
should read -- stages of Plasmodium falciparum which are homologous to --

Column 7, line 35, reads “damer sites of BcI/I and BamHI were introduced into the”
should read -- damer sites of Bc/I and BamHI were introduced into the --

Column 7, line 39, reads “NO: 1) was introduced near the BcII and BamHI linkage site”
should read -- NO: 1) was introduced near the Bc/I and BamHI linkage site --

Column 7, line 54, reads “BcII and BamHI and treated with equal volume of phenol,”
should read -- Bc/I and BamHI and treated with equal volume of phenol, --

Column 7, line 55, reads “followed by centrifugation at 12000 rpm for 5 mm. The”
should read -- followed by centrifugation at 12000 rpm for 5 min. The --

Column 7, line 59, reads “c) The digested product was ligated to vector VR1012”
should read -- c) The digested product was ligated to vector VR1 012 --

Column 7, line 60, reads “(Vical Inc.) which had been digested with same enzymes, and”
should read -- (Vical Inc.) which had been digested with the same enzymes, and --

Column 7, line 67, reads “vector VR1012.”
should read -- vector VR1 012 --

Column 8, line 4, reads “falciparum”
should read -- falciparum --

Column 8, line 11, reads “conjunction with a Hind/III site. Briefly, for the randomiza-”
should read -- conjunction with a HindIII site. Briefly, for the randomiza- --

Column 8, line 15, reads “aliquots. One was cleaved with BC/I and Hind/III the other”
should read -- aliquots. One was cleaved with BC/I and HindIII, the other --

Column 8, line 16, reads “with BamHI and Hind/III. And the fragments from the two”
should read -- with BamHI and HindIII. And the fragments from the two --

Column 8, line 64, reads “C., 3 mm 94 C., 30 sec; 50 C., 30 sec; 72 C., 30 sec; 30”
should read -- C., 3 min 94 C., 30 sec; 50 C., 30 sec; 72 C., 30 sec; 30 --

Column 9, line 1, reads “VR1O12 which had been cleaved with EcoRV and BC/I, and”
should read -- VR1 012 which had been cleaved with EcoRV and Bc/I, and --

Column 9, line 19, reads “Chimeric Genes of Plasmodium falciparum”
should read -- Chimeric Genes of Plasmodium falciparum --

Column 9, line 28, reads “The PCR product was cleaved with bc/I and BamHI and”
should read -- The PCR product was cleaved with Bc/I and BamHI and --

Column 10, line 13, reads “the gel for at least 20 mm with horizontally shaking slowly.”
should read -- the gel for at least 20 min with horizontal shaking slowly. --

Column 10, line 54, reads “from OD250/OD280 measured with DU7O ultraviolet spectro-”
should read -- from OD250/OD280 measured with DU70 ultraviolet spectro --

Column 11, line 26, reads “and 100 ul of the diluted antisera with each concentration in”
should read -- and 110 ul of the diluted antisera with each concentration in --

Column 12, lines 1-2, read “1) Recognition of Native Proteins of Plasmodium falcipar-
fum”
should read --1) Recognition of Native Proteins of Plasmodium par-
um --

Column 12, lines 3-4, read “a) Blood cells with erythrocytic stage Plasmodium falciparum 3D7 (or FCC1) (with an infection rate of about 2%)”

should read -- a) Blood cells with erythrocytic stage Plasmodium falciparum 3D7 (or FCC1) (with an infection rate of about 2%) --

Column 12, line 33, reads “native proteins of Plasmodium falciparum in above section”
should read -- native proteins of Plasmodium falciparum in above section --

Column 12, line 38, reads “Plasmodium falciparum by the antisera generated by the”
should read -- Plasmodium falciparum by the antisera generated by the --

Column 12, line 42, reads “from the Spanin-treated Plasmodium falciparum. Poly-”
should read -- from the Spanin-treated Plasmodium falciparum. Poly- --

Column 12, line 50, reads “of Plasmodium falciparum strain 3D7 was collected by cen-”
should read -- of Plasmodium falciparum strain 3D7 was collected by cen- --

Column 14, line 40, reads “of cytokine CD8, wherein positive polyepitope gene 5P312”
should read -- of cytokine CD8, wherein positive polyepitope gene SP312 --

Column 14, line 42, reads “5P352 or SP462. In contrast, negative polyepitope genes”
should read -- SP352 or SP462. In contrast, negative polyepitope genes --

Column 14, line 43, reads “behaved similar to empty vector, demonstrating that poly-”
should read -- behaved similarly to empty vector, demonstrating that poly --

Column 34, line 57, reads “a) selecting, synthesizing and cloning Into a vector a plu-” (Claim 10)
should read -- a) selecting, synthesizing and cloning into a vector a plu- --

Column 35, lines 21-22, reads “13. The method according to claim 1, wherein the ran- (Claim 13)
domly assembling of the polyepitope chimeric genes with”

should read --13. The method according to claim 1, wherein the random
assembling of the polyepitope chimeric genes with --

Column 36, lines 10-11, reads “16. The method according to claim 10, wherein the ran- (Claim 16)
domly assembling of the polyepitope chimeric genes with”

should read --16. The method according to claim 10, wherein the random
assembling of the polyepitope chimeric genes with --

Column 36, line 12, reads “different lengths in step c) is carried out simultaneously by” (Claim 16)
should read -- different lengths in step c) is carried out simultaneously by the --